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High light yield thermal neutron scintillators

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HIGH LIGHT YIELD THERMAL NEUTRON SCINTILLATORS

The invention pertains to new scintillating materials, in particular in the form of single crystals, a process to manufacture them as single crystals, and their use for to detect and/or discriminate neutron and/or gamma ray radiations.

Scintillation materials are largely used for detection of gamma-rays (or electromagnetic waves of lower energies down to 1 keV or lower, below designated as "gammas") as well as other particles like neutrons, alpha particles etc.

The scintillation mechanisms rely on a number of physical principles which essentially convert the high energy of the incoming photons or particles into light which is within or reasonably close to the visible range. Of particular interest are single crystal forms of scintillators, ie pieces that are at the scale of use constituted of one (at most a few) crystals. A single crystal (monocristal) configuration allows easier extraction of the emitted light over thick dimensions thanks to lack of internal diffusion through grain boundaries, heterogeneities and other defects that tend to be present in polycrystalline materials. A crystalline structure (in the atomic sense) is required as it determines the scintillation mechanisms: a glassy, amorphous state of matter is likely to yield different scintillation properties. The extracted light is then collected with various devices well known to the man of the art, like photomultiplier tubes, photodiodes etc. Another configuration is to still retain the crystalline structure of the material, and use it in powder form, either packed or sintered or mixed with a binder in a way that still allows light extraction. Usually, those configurations are too opaque when more than a few millimeters thick, a thickness which may not be sufficient to stop enough incoming particles or photons. Overall, whenever possible and cost effective, single crystals are preferred.

Gammas, alpha or beta particles, charged particles, or neutrons (below designated generically as "radiation") detection (is of major interest in a host of applications in nuclear medicine, fundamental physics, industral gauging, baggage scanners, oil well logging etc. In those applications, it is often desirable to discriminate neutrons from gamma rays that may also reach the detector and a scintillation detector should be able to produce different luminescence signals

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depending on the type of radiation (see G. F. Knoll, Radiation Detection and Measurements (Wiley, New York, 2000)).

Several criteria are needed to constitute a good neutron or gamma detector.

In a way well know to the man in the field, an energy spectrum of the scintillator-under-incoming-radiation-is-drawn,-whereby-events-are-represented-on-a histogram (with energies on the x-axis, the number of counts on the y-axis). In the acquisition protocol, "channels" are defined to collect the signal within a particular energy range.

Good (low) energy resolution is necessary for good energy peak identification of the incoming radiation. Energy resolution is usually determined for a given detector at a given energy as the full width at half maximum of the peak considered on an energy spectrum, divided by the energy at the centroid of the peak (see G.F Knoll, "Radiation Detection and Measurement", John Wiley and Sons, Inc, 2nd édition, p 114).

Another very important parameter is the decay time constant, which is described in particular by W.W Moses (Nucl. Instr and Meth. A336 (1993) 253). Fast decay times allow fast analyses. In general, the time spectrum of the collected signal from a scintillator under radiation (neutrons or gammas) can be fitted by a sum of exponentials characterized each by a decay time constant. The quality of a scintillator is determined essentially by the contribution of the fastest emission component. This is the number we report further in the text.

Neutron detection is often carried out with He3 tubes. This type of device has high detection efficiency, good neutron / gamma discrimination, can be made into a wide range of detector sizes, and has a long history in the field. Its weaknesses are a relatively long charge collection time and low signal levels. In a lot of cases, solid-state scintillators are preferred.

A number of scintillators used for neutron detection rely on the presence of ⁶Li in the molecule. ⁶Li is able to capture thermal neutrons and convert them into ionizing particles according to the reaction:

$${}_{3}^{6}\text{Li} + {}_{0}^{1}\text{n} \rightarrow {}_{1}^{3}\text{H} + \alpha \qquad (1)$$

The α and triton particles share a kinetic energy of 4.78 MeV. The subsequent deposition of this energy in the material can lead to detectable luminescence. However the mere presence of 6 Li does not guarantee good properties.

It is thus convenient to define a gamma / neutron discrimination indicator, the so-called γ -equivalent factor F_{γ} defined as:

$$F_{\gamma} = \frac{C_n}{C_{\gamma}} \cdot \frac{0.662}{4.78} \tag{2}$$

where C_n is the channel of the neutron peak (in the acquisition protocol) and C_{γ} the channel of the γ peak while 0.662 MeV is the energy of the γ -rays and 4.78 MeV the kinetic energy of the α and triton particles together (resulting from neutron capture).

For radiography, LiF+ZnS(Ag) screens are quite standard. The high light output (160 000 photons / neutron) is very attractive and allows good sensitivity. Its decay time is in the microsecond range.

Li glass, especially the kind enriched in ⁶Li, is another prominent neutron detector. Its light output is unfortunately very low at 6000 photons / neutron and its energy resolution high. It has the advantage of fast decay, with a decay constant at about 75 ns.

⁶Lil (lithium iodide) doped with Europium (Eu) is also a known neutron detector. The light output at about 51 000 photons / neutron is very good. The energy resolution is nonetheless extremely high, the decay constant as high as that of LiF+ZnS(Ag).

Of late, new compounds have been discovered by Bessiere et al (Scint 2003, to be published in proceedings), namely Cs_2LiYCl_6 and $Cs_2LiYCl_6:Ce^{3+}$. Both exhibit high light output (34 000 and 70 000 respectively). Their γ -equivalent factor is also very good at 0.61 and 0.66 respectively. All results on the Ce-doped version are given for a concentration of 0.1 mol%. Nonetheless, those two compounds have slow decay times (about 6-7 microseconds).

Present invention pertains to a new material showing a remarkably low decay time (often written τ). The material of the invention has as generic formula CsALiLn_(1-x)X₆: xCe³⁺, where X is either Br or I, A is Cs or Rb, Ln is Y or Gd or Lu and x is above 0.0005. The generic formula can also be written CsALiLn_(1-x)Ce_xX₆. The value x is the molar fraction of the sum of Ln and Ce. The value x is above 0.0005, but it can also be said that x is above 0.05 mol%. Both wordings are equivalent. Generally, x is less than 0.3 (= less than 30 mol%) and more generally less than 0.15.

Particularly usefull compositions are:

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- $Cs_2LiY_{(1-x)}X_6$: xCe^{3+} ,
- $CsRbLiY_{(1-x)}X_6: xCe^{3+}$.

The scintillating material of the invention can be manufactured under the form of a monocristal. In this case, its volume is generally at least 10 mm³. The material can also be used in the form of a powder (polycristalline), possibly either packed or sintered or mixed with a binder.

The compounds were grown as single crystals by the vertical Bridgman technique. Because the compounds are hygroscopic the experiments were performed on samples sealed in quartz ampoules. Other known techniques to the man in the field of crystal growths could be used such as Czochralski or Kyropoulos techniques, provided the material is reasonably protected from reaction with water and oxygen. The single crystal samples in the examples were about 10 mm³ in volume. They contain a natural abundance of ⁶Li, but they could be advantageously enriched in it (to the detriment of cost).

The properties of Cs₂LiYBr₆:Ce³⁺ are given in the Table 1 below, along with those of the scintillators already known in the field of neutron detection.

Compound	LY	R	PHD	τ
	(ph/n)	(%)	Fγ	(μs)
Cs ₂ LiYCl ₆	34 000	14.5	0.61	7.0
Cs ₂ LiXCl ₆ :0.1%Ce	70 000	5.5	0.66	6.5
Cs2LiYBr6:1%Ce	85 800 / 88 200	4.5 / 9.0	0.76 / 0.75	0.089
⁶ LiF-ZnS:Ag	160 000	_	0.45	1
⁶ LiI:Eu	51 000	40	0.86	1.2
⁶ Li glass	6 000	13-22	0.31	0.075

Table 1: Comparative properties of Cs₂LiYBr₆:Ce³⁺ and other neutron scintillators. 20

Following abbreviations were used in table 1:

- LY = Light yield
- R = energy resolution
- 25 - PHD= Puls height discrimination

 $-\tau$ = decay time

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- 0.1%Ce significate x=0.001 in the generic formula which is in fact $Cs_2LiY_{0.999}Ce_{0.001}X_6\,.$

The fast component reported for the 1mol%Ce doped version represents 39% of the emitted light, a very high number considering the overall high light yield. Pulse height discrimination (PHD), expressed as the above-mentioned F_y factor very effectively discriminates gammas from neutrons. The emission wavelength conveniently matches the maximum sensitivity of bialkali photomultiplier tubes. It has the added advantage over Cs₂LiYCl₆:0.1mol%Ce³⁺ to be extremely fast. Cs₂LiYl₆:Ce³⁺ is also of interest with the added advantage of higher detection efficiency through higher average atomic number.

The following Table 2 summarizes in more detail the properties of the compounds of the invention in the cases of gammas (as emitted by a ¹³⁷Cs source), neutrons alone or combined gammas+neutrons detection. The first two decay constants (the fastest, then second fastest) in the exponential series decomposition are reported.

	¹³⁷ Cs γ-ι	rays	Thermal neutrons		Thermal neutrons		Thermal neutrons		137Cs γ + neutrons
Compound	LY (ph/MeV)	R (%)	LY (ph/n)	Fy	R _n (%)	τ (μs)			
Cs ₂ LiYCl ₆	11 700	12.0	34 000	0.61	14.5	0.004/7.0			
Cs ₂ LiYCl ₆ : 0.1%Ce	21 600	8.0	70 000	0.66	5.5	0.004/non exp.			
Cs₂LiYBr ₆ : 1%Ce	23 600 24 700	7.0 8.5	85 800 88 200	0.76 0.75	4.5 9.0	0.089/2.5 id.			

Table 2: Main characteristics of Cs_2LiYBr_6 : Ce^{3+} under gamma, neutron and gamma + neutrons radiations (R_n = energy resolution for neutron)

It is shown in particular that materials of the invention constitute an interesting gamma detectors, less bright than the Nal:TI known in the field of gamma detection, equivalent Energy resolution but much faster decay.

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6 REVENDICATIONS

- 1- Material of the generic formula CsALiLn_(1-x)X₆: xCe³⁺, where X is either Br or I, A is Cs or Rb, Ln is Y or Gd or Lu and x is above 0.0005.
- 5 2- Material according to preceeding claim, wherein x is above 0.005.

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- 3- Material according to either of preceeding claims, wherein x is less than 0.3.
- 4- Material according to preceeding claim, wherein x is less than 0.15.
- 5- Material according to either of preceeding claims, wherein it is under the form of a monocristal.
- 6- Material according to preceeding claim, wherein the volume of the monocristal is at least 10 mm³.
- 7- Material according to either of claims 1 to 6, wherein it is under the form of a powder.
- 8- Material according to preceeding claim, wherein it is either packed or sintered or mixed with a binder.
 - 9- Material according to either of preceeding claims, wherein its formula is $Cs_2LiYX_6:xCe^{3+}$.
- 10- Use of material according to either of preceeding claims in gamma and / or neutron detection.

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ABSTRACT

The invention pertains to a scintillating material of the generic formula $CsALiLn_{(1-x)}X_6: xCe^{3+}$, where X is either Br or I, A is Cs or Rb, Ln is Y or Gd or Lu and x is above 0.0005, generally under the form of monocristal. This material has a remarkably low decay time. It can be used to detect and/or discriminate neutron and/or gamma ray radiations.

